DOCUSATE SODIUM- docusate sodium capsule, liquid filled
WALGREEN COMPANY

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Walgreens Gentle Stool Softener

Drug Facts

Active ingredient (in each liquid-filled capsule)

Docusate Sodium 100 mg

Purpose

Stool Softner laxative

Uses

- relieves of occasional constipation (irregularity)
- this product generally produces a bowel movement within 12 to 72 hours

Warnings

Ask a doctor before use if you have

- stomach pain, nausea or vomittiong
- a sudden change in bowel habits that lasts more than 2 weeks

Do not use if you are presently taking mineral oil, unless told to do so by a doctor

Stop use and ask a doctor if you

- you have rectal bleeding ot no bowel movement after using this product. These could be signs of a serious condition.
- yo need to use laxative for more than 1 week

If pregnant or breast-feeding,ask a health professional before use

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.1(800)222-1222

Directions

take with a glass of water

take by mouth. Doses may be taken as a single daily dose or in divided doses.

Adults and children 12 years and over	take 1 to 3 capsules daily.
Children 2 to under 12 years of age	1 capsule daily.
Children under 2 years of age	Ask a doctor

Other information

- Store at room temperature 20-25°C (68-77°F)
- protect from excessive humidity
- do not use this product of the safety seal under the cap is torn or missing

Inactive ingredients

FD&C Red#40, FD&C Yellow#6, gelatin,glycerin, polyethylene glycol 400, propylene glycol, povidone, purified water, sorbitol solution, titanium dioxide.

Questions or comments?

1-888-333-9792

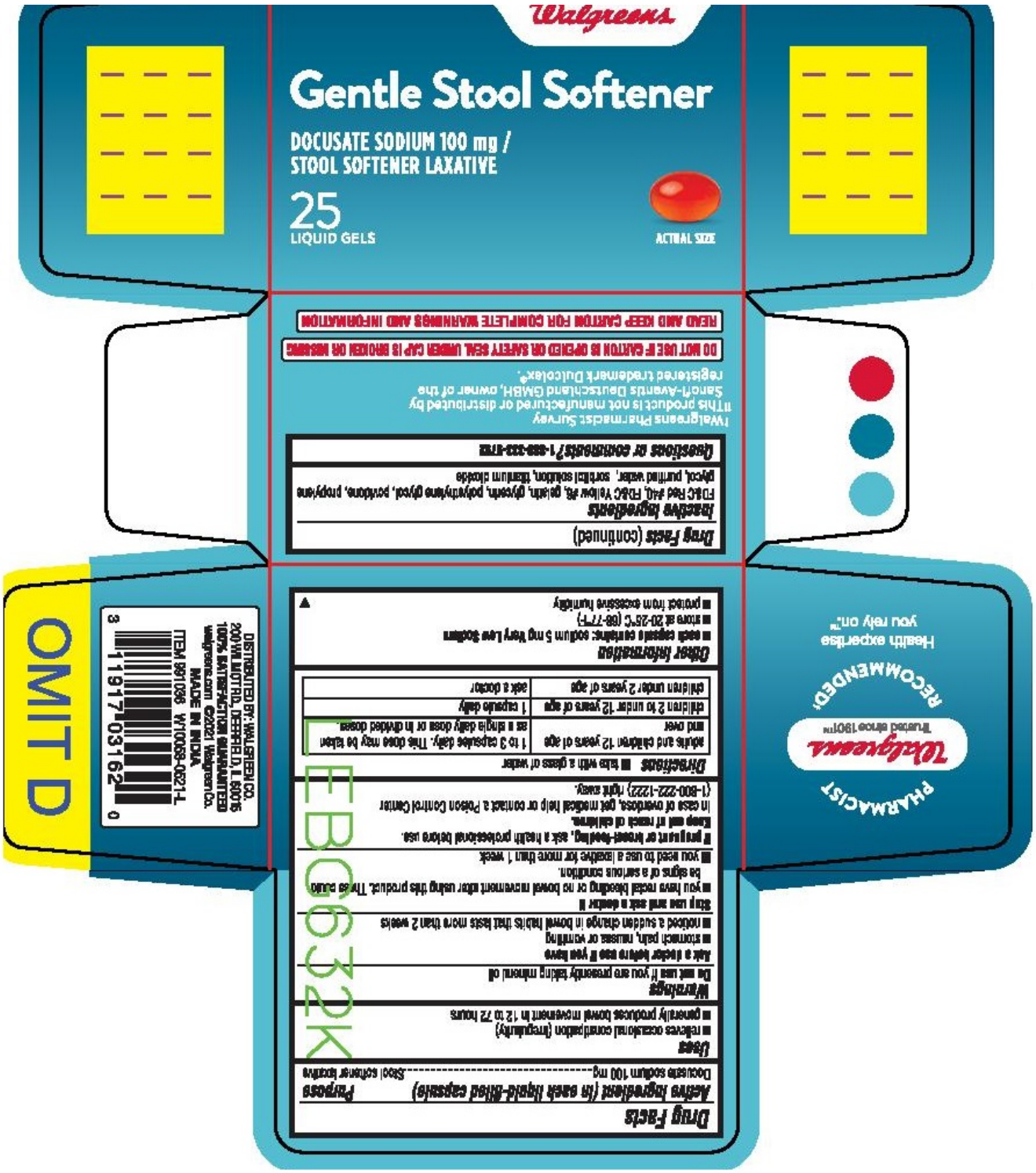
PRINCIPAL DISPLAY PANEL

Compare to Dulcolax ® Stool Softener active ingredient *

GENTLE STOOL SOFTENER

DOCUSATE SODIUM 100 mg / STOOL SOFTENER LAXATIVE

25 LIQUID GELS



DOCUSATE SODIUM

docusate sodium capsule, liquid filled

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-8991
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DOCUSATE SODIUM (UNII: F05Q2T2JA0) (DOCUSATE - UNII:M7P27195AG)	DOCUSATE SODIUM	100 mg

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
SORBITOL (UNII: 506T60A25R)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POVIDONE K30 (UNII: U725QWY32X)	

Product Characteristics

Color	red	**Score**	no score
Shape	OVAL	**Size**	12mm
Flavor		**Imprint Code**	125
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-8991-25	1 in 1 PACKAGE	12/22/2020	
1		25 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part334	12/22/2020	

Labeler - WALGREEN COMPANY (008965063)

Revised: 11/2021 WALGREEN COMPANY